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PACRAT: A MACHINE BRAIN

PART ONE

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INTRODUCTION

The vertebrate brain, and in particular the mammalian brain, consists of:

1. a neural net, the cortex, that responds differentially to the exterior universe,
2. motor programs hardwired in the basal ganglia,
3. a throttle, the thalamic reticular nucleus, that can stop the flow of signal energy from the sensory neurons to the neocortex, and also the flow of motor programs from the basal ganglia to the motor cortex,
4. an averager, the cerebellum, that keeps running averages of the signals sent to the muscles and incoming signal information.

Associated with this brain is a mind. The association is not part of the universe of experience.

BACKGROUND

All of the neurons in the central nervous system are equal; but some are more equal than others. What are the essentials? We need a nerve net that responds differentially to the universe. Response to oriented line segments and the orthogonal movement of these segments have been identified (Hubel, 1988). We need a set of motor programs provided by the genes and we nominate the basal ganglia to provide them. We need an interruption mechanism to disconnect the motor program from the motor neuron. The thalamic reticular nucleus is a likely candidate. Lastly we need modifiable synapses. The Hebbian hypothesis currently serves. If a synapse participates in firing a neuron, it is strengthened.

By motor program we mean a hardwired neural net in which neuron A excites neuron B and 10 msec later neuron C. Neuron B and C are connected to motor neurons so that one muscle fiber contracts 10 msec before the other and a delicate movement is produced. The essential quality of the motor program is that it is hardwired. The brain does not "recall" the motor program; the signal energy triggers it. Also, we speak of two neurons rather than 2000.

We cannot see the mind as object. Nothing rational can be said of the mind. It is the subject for which the world is object.

The guiding question for this endeavor is: Will this notion advance the design of an electronic brain?

The Unknowable

We look at consciousness from the outside, we see an animal that is alert. When I look inside, I am aware. I need not think; to be aware is enough. The brain and the mind may be the same thing, may be different aspects of the same thing, may be totally different things. We do not know; we cannot know. Man cannot know the relationship of 'alert' to 'aware.'

Mind-Body

The mind is aware; the brain is alert.

In the beginning, the mind watched; the brain thought and the mind was aware of the thought. Awareness is of the mind. This quality of awareness is a mystery for which we have no explanation.

I am aware.
You are alert.
He exhibits intelligent behavior.

This is the box in which we live. Is there a way out? We doubt it. Although we gladly admit all vertebrates as potentially aware--certainly all mammals (Kissin, 1986:81) and probably octopi and squid--we have never found an experiment that gives any hint of testing for awareness. Turing's test tells us nothing; he speaks only of intelligence.

All brain activities reduce to a pattern of connection. All mental events reduce to awareness of neural activation.

Any animal that is alert may be aware.
A machine may be alert.
Any animal that has plastic synapses may learn.
A machine may have plastic synapses.

The metalanguage differs from rational language. We may not use rational language to talk about minds. Minds only experience rationalism. We may use rational language to talk about brains.

We are under no direction to speak only rationally.

Awareness

Consciousness of the outer world occurs at points. One neuron (or one glomerulus) in the lateral geniculate nucleus, rising to awareness, is enough. The mind is aware of the activity of single neurons. We readily sense a point source of light that excites, at most, only a few neurons. The mind is not equally aware of all neurons. Some neurons rank high, whereas other neurons are silent.

The Homunculus

The homunculus is a little man who sits in the middle of the brain watching a television screen and pushing buttons. The homunculus stores memories, retrieves them, and compares them with present experience. We exorcise the homunculus by postulating a passive brain. The mind watches the signal energy coursing through the brain, but the brain knows nothing. It neither stores, nor retrieves, nor compares. It is wholly passive. We need a valve to shut off the flow of motor output temporarily. The thalamic reticular nucleus is in the right place and is capable of doing it. "What function the reticular nucleus really effects is also largely a matter of conjecture" (Jones, 1985:819).

The Passive Brain

We see a brain composed of neurons and the neurons as generators of electrical pulses. Signal energy courses through this passive brain. There is no need for rigor here. We say most of them generate pulses, with a minority that instead deals in graded potentials. We know enough about the brain to imagine how it works, but nowhere near enough to do anything but guess.

We may view the brain as electronic circuitry. We can move the abstract circuitry from one substrate to another, from organic structure to silicon.

Thought

E. V. Evarts quoted Walle Nauta as having said "What is a thought except a movement that is not connected to a motor neuron" (Ciba Foundation, 1984:272).

Schneider thought Nauta might be stretching the point slightly; we think not (Schneider, 1987:7). We think it is a simple, straightforward statement that needs no qualification. The manner of the disconnection is what we must examine. Just how do we disconnect the motor program from the motor neuron in a passive brain? The disconnection is not physical, of course...no axon dangles in space. It must be done electrically, by inhibition. The only nucleus that is in a position to do this is the reticular nucleus of the thalamus. It is in position to interrupt the flow from the basal ganglia to the motor cortex.

Thought has two faces. One is the mechanical action of the brain; the other is the awareness of this action. This awareness is that which philosophers have traditionally called thought. It is in this talk that we find mental images and logical forms. We have no need of these notions. The style of thinking that interests us is that of Nauta. We would like to say it is the brain that thinks, with the mind as spectator only. The mind is aware of thinking as it is of pain and happiness.

Thinking follows when the reticular nucleus of the thalamus inhibits sensory input. The signal energy that has already passed through is then free to vibrate and oscillate. We present thinking as a mindless activity of the brain of which the mind is only aware. This we see as the best way to approach the design of an electronic brain. Extended thought is a luxury of a dominant species at the top of the food chain, an aberration, deviant. Thought has marginal utility.

THE PROGRAM: PART ONE

The program, Pacrat, contains a brain, an organism, and an environment in which the organism moves about as directed by the brain. The program splits logically into three parts. The first defines the brain, the organism, and the universe. The second exercises the brain. The third moves Pacrat and evaluates the response of the universe. A subsidiary program, running on a workstation, provides the graphic output that allows us to judge the actions of Pacrat. Here we describe the first part of the main program.

COMMON

We designed the program about named common. The common blocks completely define the condition of the brain, the stance of the organism, and the situation of the environment. The program manipulates the elements in the named common. An appendix sets forth their structure.

MAIN

The main program calls a subroutine, SETUP, that builds a brain and an environment. It then calls CYCLE in a do loop. CYCLE steps through the thinking, movement, and the reaction of the environment.

```
program pacrat
include 'brain'
chkout=.false.
call setup
call lister
do 20 j=1,kbgend
do 10 k=1,kltend
call cycle
```

```

    call chflag
    if(mod(icnt,ksnap).eq.0) call hotout
10  continue
20  continue
    call closer
    call exit
    end

    subroutine cycle
    include 'brain'
    call updacl
    call ras
    call update
    if(mod(icnt,10).eq.0) then
c    call weight
    end if
    if(.false.) then
c    if(mod(icnt,100).eq.0) then
    write(*,*) 'icnt',icnt
    call growth
    call recept
    call nusyns
    end if
    if(mod(icnt,10).eq.0) then
    call mover
    call univrs
    if((icnt.ge.0).and.(icnt.le.100))then
    call actvty
    end if
    end if
    return
    end

```

SETUP

The first call of the main program is to SETUP. SETUP opens all the working files. The working routines open the checkout files where the action occurs.

brain.nms	contains the names of the brain centers
brain.cnx	specifies the efferent connections and the scheme of connection
walls	the parameters of the universe
trail	history of Pacrat's progress
brain.ckr	
brain.strt	parameters for a given run

brain.snap	the entire contents of the brain for a hot start
------------	---

Following the opening of files, we read in a record from brain.strt containing starting parameters. The following variables define the run:

chkout	a checkout flag
hotstr	true calls for a hot start; we read in a brain snapshot
ksnap	an interval (in cycles) between between brain snapshots
kaix	a starter for the random number generator
kbgend	terminator for the outer loop in the main program
kltend	terminator for the inner loop

Next, we call RANDIT(kaix) to initialize the random number generator. We use the random numbers to avoid the artificial regularity inherent in the serial makeup of Fortran do loops. If two or more synapses are in the same condition, we add a small random increment so that neurons will be selected in an arbitrary order.

Next we read in information about desired check out using the namelist method. The namelist contains:

flags (100)	(logical) these flags control the check out listings
lstnr(10)	the number of entries in nuclst
nuclst(100)	(character*30) the names of the nuclei that we want to examine
kbndls(3)	these numbers control the do loop that lists the status of neurons

At this point we decide, depending on the truth value of hotstr, whether to make a hot start by calling HOT. The subroutine HOT will read in the contents of the brain as it was at the last snapshot. Alternatively we call COLD to build a brain from parametric information.

```

subroutine setup
include 'brain'
namelist/snaps/ flags,lstnr,nuclst,kbndls

```

```

c
c all working files are opened with the exception of
c brain.snap that will be opened in subroutine hot if reading
c or writing a snapshot of the brain is called for.
c

```

```

open(1,file='brain.nms',status='old')
open(2,file='brain.cnx',status='old')

```

```

open(3,file='walls',status='old')
open(4,file='trail')
open(8,file='brain.ckr')
open(9,file='brain.strt',status='old')
c
c 11,12,13, and 18 are check-out files.
c
open(20,file='radii',status='old')
read(9,10) chkout,hotstr,ksnap,kaix,kbgend,kltend
write(*,*) chkout,hotstr,ksnap,kaix,kbgend,kltend
10 format(2I5,4I10)
c
c  chkout  a checkout flag
c  hotstr  if true a hot start is called for, a brain
c          snapshot will be read in.
c  ksnap   the interval at which brain snapshots will be taken.
c  kaix    a starter for the random number generator.
c  kbgend  outer loop in main
c  kltend  inner loop in main
c
c the random number generator is initialized.
c
aix=randit(kaix)
read(9,snaps,end=35)
35 continue
c
c a selection of flags and bounds are read in. These are used
c to print out states of the brain during program execution.
c
c  flags(100)  logical      these are flags used to direct
c                  monitoring.
c  flags(1)    In subroutine chflag, if true sends control to
c                  chekpr.
c
c  lstnr(10)
c  lstnr(1)    number of entries in nuclst.
c
c  nuclst(100) character*30
c              a list on nuclei to be printed out as they are called for.
c
c  kbndls(3)
c  kbndls(1), kbndls(2), and kbndls(3) form the parameters
c  of a do loop in chekr.
c
c
c select either a hot or a cold start.
c
if(hotstr) then
call hot
else
call cold
end if

```

```

c
c initialize hubel
c
c      call hublin
c
c initialize visual
c
c      call visiin
c
c initialize motr
c
c      call motrin
c      return
c      end

      function randu(x)
      save
      iy=iy*65539
      if(iy.lt.0) iy=iy+2147483647+1
      randu=iy*.4656613e-9
      return
      entry randin (ix)
      iy=ix/2
      iy=iy*2+1
      randin =0.0
      return
      end

      function gauss(sigma,amean)
      dimension f(10,10)
      save
      a=0.0
      do 50 i=1,12
50  a=a+randu(z)
      gauss=(a-6.0)*sigma+amean
      return
      entry randit(ix)
      call randin(ix)
      do 60 j=1,10
      do 60 k=1,10
60  f(j,k)=randu(z)
      randit=0.0
      return
      entry rand(x)
      j=10.0*randu(x)+1.0
      k=10.0*randu(x)+1.0
      swap=f(j,k)
      f(j,k)=randu(x)
      rand=swap
      return
      end

```

HOT

This routine reads in the contents of the named commons from a snapshot file that we established in a previous run by a call to HOTOUT. This restarts Pacrat as if there had been no intermission.

HOTOUT

HOTOUT opens a file called brain.snap and dumps the entire contents of the named commons. This large file contains all the numbers and logical variables that define the brain, the organism, and the environment. These are the synaptic connections with their facilitation and conditions; a history of the neurons, and all the numbers that make for a simulation; the description and location of the organism; and a description of the universe.

subroutine hot

```
c
c when we rewrite this for new common, watch out for
c line length.
c
  include 'brain'
  open(10,file='brain.snap',status='old',
1    form='unformatted')
  read(10) a,b,centrs,nuclei,nrctr,nrnu,centrs,
1    nuclei,kcnucs,kruse,krnet,krgr,ksens,nucnet,
2    nucgro,kcensz,knucsz,knucwd,stinc,knucs,kside
  read(10) ksynps
  read(10) potent
  read(10) synred
  read(10) ksynct,histry
  read(10) ktime
  read(10) alertd,runavg,thresh,recptv,grfctr,synare,grinc,
4    icnt,reduce,frustr,frlvla,frlvlb,rewavg,recovr,
5    thrup,thrdwn
  read(10) chkout,hotstr,ksnap,knrrad,radii,body,nrwall,
6    awalls,avectr,amoton,bmoton,amovr,phase,radi,
7    snpass,ampass,dist,dxm
  close (10)
  return
  entry hotout
  open(10,file='brain.snap',form='unformatted')
  write(10) a,b,centrs,nuclei,nrctr,nrnu,centrs,
1    nuclei,kcnucs,kruse,krnet,krgr,ksens,nucnet,
2    nucgro,kcensz,knucsz,knucwd,stinc,knucs,kside
  write(10) ksynps
  write(10) potent
  write(10) synred
  write(10) ksynct,histry
  write(10) ktime
  write(10) alertd,runavg,thresh,recptv,grfctr,synare,grinc,
4    icnt,reduce,frustr,frlvla,frlvlb,rewavg,recovr,
5    thrup,thrdwn
  write(10) chkout,hotstr,ksnap,knrrad,radii,body,nrwall,
6    awalls,avectr,amoton,bmoton,amovr,phase,radi,
```

```

7      snpass,ampass,dist,dxm
      close (10)
      return
      end

```

COLD

This routine schedules all the routines that build the brain. It calls in turn INIT, INITX, NAMER, NET, and CONNEX. After these calls, the routine returns control to SETUP.

```

      subroutine cold
c
c This routine creates a brain by calling the following
c subroutines.
c
      include 'brain'
      call init
      call initx
      call namer
      call net
      call connex
      return
      end

```

INIT

INIT establishes several parameters, and resets the cycle count, icnt. Then it defines certain bounding values:

memsiz	an upper bound on the number of neurons
kcensz	an upper bound on the number of centers
knucsz	an upper bound on the number of nuclei
kside	the number of neurons on a side of a nucleus
knucwd	an upper bound on the number of nuclei to which this nucleus may have efferent connections, either genetic or epigenetic

The routine establishes parameters about the dynamic action of the brain:

thrup	increment of threshold hyperpolarization
thrdwn	increment of threshold hypopolarization
reduce	decrement of synaptic reserve

recovr	increment of synaptic reserve
stinc	synaptic strength at genetic establishment
grinc	synaptic strength at epigenetic growth
peaked	(a skeleton)
frlvla	upper level of frustration
frlvlb	lower level of frustration
dxm	a step size for movement of the organism
phase	original value for sinusoidal movement

The routine returns control to COLD.

subroutine init

include 'brain'

c

c Certain basic parameters are declared.

c	icnt	the cycle counter
c	kcensz	an upper limit on the number of brain centers
c	knucsz	an upper limit on the number of nuclei
c	kside	the number of neurons on one side of a nucleus.
c		the nuclei are square.
c	knucwd	an upper limit on the number of nuclei that the
c		neurons in this nucleus can be efferent upon or
c		grow to.
c	thrup	hyperpolarization increment
c	thrdwn	hypopolarization decrement
c	reduce	decrement in synaptic availability
c	recovr	increment in synaptic availability
c	stinc	genetic synaptic potentiation
c	grinc	epigenetic synaptic potentiation
c	frlvla	upper check on frustration (frustr)
c	frlvlb	lower check on frustration
c	dxm	a step size for movement of organism
c	phase	an initial value for sinusoidal movement

c

```

icnt=0
kcensz=100
knucsz=300
kside=16
knucwd=20
thrup=0.003
thrdwn=0.001
reduce=0.003
recovr=0.001
stinc=0.2

```

```

grinc=0.01
frivla=0.9
frivlb=0.1
dxm=0.05
phase=0.0
return
end

```

INITX

This routine resets all the arrays that are to contain the dynamic history of the brain. These are: kruse, krnet, krgro, nucnet, nucgro, ksynct, thresh, and histry. We initialize runavg at 0.5 and reset alertd, ksens, and snpaso. Then we read in a description of the universe into awalls from the file walls, and place the starting position of the organism in amovr. We place the position of the head of the organism in avectr and amoton, and reset the working array, bmoton.

Next we read in a description of the organism from the file radii and place it in the array, radii.

We return control to COLD.

```

subroutine initx
include 'brain'
do 5 j=1, knucsz
kruse(j)=0
krnet(j)=0
krgro(j)=0
do 5 k=1, knucwd
do 5 l=1, 4
nucnet(j,k,l)=0
5 nucgro(j,k,l)=0
do 10 l=1, knucsz
do 10 k=1, kside
do 10 j=1, kside
ksynct(j,k,l)=0
thresh(j,k,l)=0.0
10 continue
do 110 l=1, knucsz
do 110 k=1, kside
do 110 j=1, kside
do 105 i=j, 10
histry(i,j,k,l)=0.0
105 continue
110 continue
do 120 i=1, knucsz
runavg(i)=0.5
alertd(i)=0.0
120 continue
read(3,25) nrwall
25 format(i5)
do 30 k=1, nrwall
read(3,27) ((awalls(i,j,k), i=1,3), j=1,2)
27 format(6f10.5)

```

```

30 continue
   close(3)
   do 45 k=1,100
   do 44 j=1,3
   amovr(j,k)=0.0
44 continue
c
c set following for initial displacement of pacrat.
c
   amovr(1,k)=7.0
c   amovr(3,k)=5.0
45 continue
   do 46 j=1,3
   avectr(j)=0.0
   do 43 i=1,3
   amoton(i,j)=0.0
   if(i.eq.j) amoton(i,j)=1.0
43 continue
46 continue
c
c special coding to start at a 45 degree angle.
c
   arg=3.141593/4.0
   amoton(1,1)=cos(arg)
   amoton(3,3)=cos(arg)
   amoton(1,3)=-sin(arg)
   amoton(3,1)=sin(arg)
c
   do 90 l=1,100
   do 90 k=1,3
   do 90 j=1,3
   bmoton(j,k,l)=0.0
   if(j.eq.k) bmoton(j,k,l)=1.0
90 continue
   read(20,47)knrrad
47 format(i5)
   do 60 k=1,knrrad
   read(20,48) (radii(1,j,k),j=1,3,2)
48 format(2f10.5)
   radii(1,3,k)=radii(1,3,k)/3.0
   radii(1,2,k)=0.0
   radii(2,3,k)=-radii(1,3,k)
   do 50 j=1,2
   radii(2,j,k)=radii(1,j,k)
50 continue
60 continue
   return
   end

```


NAMER

This routine reads in the names of the centers and nuclei that are to form the brain. We verify that no name duplicates a previously defined name and store the center in `centrs` and the nuclei in `nuclei`. We place the count of centers in `nrctr` and the count of the nuclei in `nrnucs`, then return control to `COLD`.

```
      subroutine namer
      include 'brain'
c
c it is assumed that all the names of centers and of nuclei
c will be read in. there will be a center and a nucleus on
c each card. the centers will appear in groups, and no nucleus
c will be repeated.
c
c why the following appears is not known
c
      kadhoc=77
c
      kctr=1
      knucs=1
      read(1,20,end=100) a,b
      centrs(kctr)=a
      nuclei(knucs)=b
      kcnuks(kctr,1)=knucs
c
c this is the main loop.
c
      10 continue
      read(1,20,end=100) a,b
      20 format(2a30)
c
c check that the nucleus is unique
c
      call ident(b,jn,kn)
      if((jn.ne.0).and.(kn.ne.0)) then
      write(13,35) a,b
      35 format(1h,'dupe ',2(a30,2x))
      end if
c
c is this a new center
c
      if(lge(centrs(kctr),a).and.lle(centrs(kctr),a)) then
      knucs=knucs+1
      if(knucs.gt.knucsz) then
      write(13,45) knucs,knucsz
      45 format(1h,'exceeded knucsz in namer',2i10)
      call exit
      end if
      nuclei(knucs)=b
      else
c
c it is.
```

```

c
    kcnuks(kctr,2)=knucs
    kctr=kctr+1
    knucs=knucs+1
    if((knucs.gt.knucsz).or.(kctr.gt.kcensz)) then
    write(13,55) kctr,kcensz,knucs,knucsz
55  format(1h,'exceeded kcensz or knucsz in namer',4i10)
    call exit
    end if
    kcnuks(kctr,1)=knucs
    centrs(kctr)=a
    nuclei(knucs)=b
    end if
c
c nrctr and nrnucs are to contain the total of centers and nuclei.
c
    kcnuks(kctr,2)=knucs
    go to 10
c
c continue the main loop.
c
100 continue
    nrctr=kctr
    nrnucs=knucs
    do 120 k=1,nrctr
    write(18,110) k,kcnuks(k,1),kcnuks(k,2)
110 format(3i5)
120 continue
    return
    end

```

IDENT

This routine returns the center, j, and the nucleus, k, of the name, a. If the name does not appear in the list of nuclei, we reset j and k.

```

    subroutine ident(a,j,k)
    include 'brain'
c
c this routine depends upon nuclei appearing only once.
c
    do 40 ja=1,nrctr
    do 30 ka=kcnuks(ja,1),kcnuks(ja,2)
    if((lle(a,nuclei(ka)).and.lge(a,nuclei(ka))))then
    j=ja
    k=ka
    return
    end if
30 continue
40 continue
    j=0

```

```

k=0
return
end

```

NET

NET reads in a description of the neural connections that define the brain: lx, ly, lw, lz, a, and b, where a is the name of the nucleus in question and b is the name of the nucleus to which it is afferent. Of course, each nucleus may be afferent to more than one other nucleus, including itself.

We check a and b to see that they are present in the list of nuclei in nuclei. We do this check in IDENT and then record the following distinctions:

lx=1	this is a reservation of space
lx=2	genetic instructions follow
lx=3	epigenetic instructions follow

If lx=1, then ly is the square root of the number of neurons to be reserved for this nucleus. If lx=2 or 3, then ly is the number of efferent synapses for each neuron in this nucleus. If ly=10, then the first digit identifies the scheme of connection (other than topological).

If lw=3, this is a sensory nucleus. If lw=4, this is a motor nucleus.

We use lz to make a distinction between ipsilateral and contralateral synapses, and between excitatory and inhibitory connections.

lz=1	synapses are ipsilateral and excitatory
lz=2	synapses are ipsilateral and inhibitory
lz=3	synapses are contralateral and excitatory
lz=4	synapses are contralateral and inhibitory

The call to IDENT returned a value, kn, that is the identifying number of this nucleus. A second call to IDENT returns the value, kna, that identifies the afferent nucleus. Kn and kna index the nucleus name in nuclei and the position of this nucleus in the brain arrays. If kruse(kn)=0, we set it to 1 to show that this nucleus is active. If lw=3, we set kruse(kn) to 3 to identify it as a sensory nucleus. If lw=4, we set kruse(kn) to 4 to identify it as a motor nucleus.

If lx=2, we increment krnet(kn) and check it against knucwd for overflow. We place the values of kna, ly, lw, and lz in nucnet(kn,krnet(kn),__) where __ is 1, 2, 3, or 4.

If lx=3, we increment krgro(kn) and check it as we did for lx=2, and place the values of kna, ly, lw, and lz in nucgro.

The routine returns control to COLD.

```

subroutine net
include 'brain'

```

c

```

c this routine reads in the parameters of the neural net.
c
c if lx=1 this is a reservation only (a skeleton).
c   =2 synaptic instructions
c   =3 growth instructions
c
c if lx.eq.1 then
c   ly is the square root of the neurons to be reserved.
c   for 2 or 3, ly is the number of efferent synapses.
c if lx.eq.2 or lx.eq.3 then
c   ly is the number of efferent synapses with straight
c   (randomized possibly) topological connection.
c if ly.gt.10 then
c   the first digit codes the type of synaptic
c   distribution as in Hubel.
c
c if lw=3 this is a sensory nucleus
c   =4 this is a motor nucleus
c
c lz=1 synapses are excitatory
c   =2 synapses are inhibitory
c   =3 contralateral excitatory
c   =4 contralateral inhibitory
c
10 continue
   read(2,15,end=300) lx,ly,lw,lz,a,b
15 format(4i5,2a30)
c
c here we get the number of the efferent nucleus
c
   call ident(a,jn,kn)
   if((jn.eq.0).or.(kn.eq.0))then
   write(13,25) a
25 format(1h,'in net, ',a30,' not found in names')
   end if
   if(kruse(kn).eq.0) kruse(kn)=1
c
c the reservation of the square of ly is a skeleton and is
c negated here.
c
   if(lx.eq.1) go to 10
   if(lw.eq.3) kruse(kn)=3
   if(lw.eq.4) kruse(kn)=4
c
c here we get the number of the afferent nucleus
c
   call ident(b,jna,kna)
   if((jna.eq.0).or.(kna.eq.0))then
   write(13,25) b
   write(*,*) 'in net failed to find',b
   end if
   if(lx.eq.2) then

```

```

krnet(kn)=krnet(kn)+1
if(krnet(kn).gt.knucwd) krnet(kn)=knucwd
nucnet(kn,krnet(kn),1)=kna
nucnet(kn,krnet(kn),2)=ly
nucnet(kn,krnet(kn),3)=lw
nucnet(kn,krnet(kn),4)=lz
go to 10
end if
if(lx.eq.3) then
krgro(kn)=krgro(kn)+1
if(krgro(kn).gt.knucwd) krgro(kn)=knucwd
nucgro(kn,krgro(kn),1)=kna
nucgro(kn,krgro(kn),2)=ly
nucgro(kn,krgro(kn),3)=lw
nucgro(kn,krgro(kn),4)=lz
go to 10
end if
go to 10
300 continue
return
end

```

CONNEX

We evaluate ly for class of connection and make a call either to TOPO for topological efferents or to HUBEL for filtered efferents. Following these calls, we call SYNARM for each neuron to normalize the afferent synapses if it should be necessary.

The routine returns control to COLD.

Connections in the Cortex

The function of the cortex is to accept signal energy, primarily from the thalamus, and provide multiple paths to the caudate nucleus. We may think of these multiple paths as providing filtration, if we like. The signal energy turns on neurons in the cortex according to patterns, patterns that are not intuitive, not what we would expect. The incoming visual energy has already arranged itself in the retina in center-on and center-off patterns. These patterns consist first of a field, a region in visual space in which light will have an effect on the output neuron. The field has a center and a surround. The center excites and the surround inhibits.

On entering the koniocortex these center-surround signals arrange themselves in line segments. The next arrangement involves motion perpendicular to a line segment. The first represents a pattern in space; the second, a pattern in time of the spatial pattern.

The more complex patterns can only be simple combinations in space and time of the patterns in a previous column or area.

Since little may be said about these patterns beyond the very first ones, we will pretend that they are repetitions of the primary patterns. It is our most fundamental belief that they must be simple. We intend to add routines that will discriminate moving lines and then lines moving from or to the center. These will provide the neural analog of "things moving toward" and "things moving away."

```

subroutine connex
include 'brain'
c
c the kind of connection is looked at. it is either a filter or
c topographic at present.
c
do 70 i=1,nrnucs
if(kruse(i).eq.0) go to 70
if(krnet(i).eq.0) go to 70
do 60 j=1,krnet(i)
if(nucnet(i,j,2).ge.10) then
call hubel(i,j)
else
call topo(i,j)
end if
60 continue
70 continue
c
c the afferent synapses are normalized.
c
do 80 l=1,nrnucs
do 80 k=1,kside
do 80 j=1,kside
80 call synnrm(j,k,l)
return
end

```

TOPO(lf,lnr)

This routine connects each neuron in the lfth nucleus to the equivalent neuron in the lnrt nucleus in its table of genetic connection. If there is to be more than one synapse per neuron, there is provision for random selection of neighboring neurons. A call to INSERT establishes the synapse. The routine then returns control to CONNEX.

We feel that this is a basic pattern of connection but there is always and accompanying filtration. The preservation of topology is a strong parameter in the genetic organization of a brain.

```

subroutine topo(lf,lnr)
include 'brain'
c
c lf is nucleus from, lnr is index of nucleus to
c
c
apot=stinc
ltime=1
lt=nucnet(lf,lnr,1)
ly=nucnet(lf,lnr,2)
lw=nucnet(lf,lnr,3)
lz=nucnet(lf,lnr,4)
c
c lw identifies the target nucleus as possibly sensory
c or motor.
c ly is the number of synapses from to going.

```

c lz indicates type of synapse and whether ipsilateral
 c or contralateral.
 c

```

    delta=1.0
    sigma=delta/2.36
    krdm=ly*delta**2
    if(krdm.lt.1) krdm=1
    do 40 kf=1,kside
      akf=kf
      do 30 jf=1,kside
        ajf=jf
        do 24 kqr=1,krdm
          27 continue
          kt=gauss(sigma,akf)
          jt=gauss(sigma,ajf)
          if((kt.lt.1).or.(kt.gt.kside)) go to 27
          if((jt.lt.1).or.(jt.gt.kside)) go to 27
          if(kqr.eq.1) then
            jt=jf
            kt=kf
          end if
          call insert(jf,kf,lj,jt,kt,lt,apot,ltime,lz)
          if(jt*kt*lt.eq.0) go to 27
        24 continue
      30 continue
    40 continue
    return
  end

```

HUBEL(lf,lnr)

This, like TOPO, is a genetic connection routine. Here we attempt a more significant distribution of efferent synapses. We use Hubel's directed line segments as our choice to add some realism to our intercortical connections. A call to INSERT establishes the synapse.

This routine effects a complex mapping from the efferent nucleus to the cortex. Each cortical neuron responds to a line segment. The segment itself has an inhibiting zone on both sides. We do not see this inhibiting effect occurring here, but in the previous stage. As we pass through the cortex we come to ever more complex cells, but the complexity is not in the immediate cell but in the previous cells. The inhibiting borders are a natural outcome of what came before: the center on, surround off cell.

The routine returns control to CONNEX.

```

subroutine hubel(lf,lnr)
include 'brain'

```

```

c
c lf is nucleus from, lnr is index of nucleus to
c
  dimension ktrial(16,16,16,2)
  apot=stinc
  !time=1
  lt=nucnet(lf,lnr,1)

```

```

ly=nucnet(lf,lnr,2)
lw=nucnet(lf,lnr,3)
c   nucnet(lf,lnr,4)=mod(nucnet(lf,lnr,4),10)  why this here?
lz=nucnet(lf,lnr,4)
do 150 jta=1,kside
do 140 kta=1,kside
do 130 ii=1,kside
jt=jta
kt=kta
lt=nucnet(lf,lnr,1)
jf=kttrial(ii,kt,jt,1)
kf=kttrial(ii,kt,jt,2)
call insert(jf,kf,lf,jt,kt,lt,apot,ltime,lz)
if(jt*kt*lt.eq.0) go to 130
130 continue
140 continue
150 continue
    return
    entry hublin
    aone=0.1989124
    atwo=0.414214
    athree=0.668179
    do 10 j=1,kside
    kttrial(j,1,1,1)=1
    kttrial(j,1,2,1)=(kside-j)*aone+1.5
    kttrial(j,1,3,1)=(kside-j)*atwo+1.5
    kttrial(j,1,4,1)=(kside-j)*athree+1.5
    kttrial(j,1,5,1)=kside-j+1
10  continue
    do 20 l=1,5
    do 20 k=2,kside
    do 20 j=1,kside
    w=kttrial(j,k-1,1,1)+1
    if(w.gt.kside) w=w-kside
    kttrial(j,k,1,1)=w
20  continue
    do 30 l=1,5
    do 30 k=1,kside
    do 30 j=1,kside
    kttrial(j,k,1,2)=j
30  continue
    do 40 l=6,9
    la=10-l
    do 40 k=1,kside
    do 40 j=1,kside
    kttrial(j,k,1,1)=kttrial(j,k,1,2)
    kttrial(j,k,1,2)=kside-kttrial(kside-j+1,k,la,1)+1
40  continue
    do 50 l=10,13
    la=18-l
    do 50 k=1,kside
    do 50 j=1,kside

```



```

    ktrial(j,k,l,1)=ktrial(j,k,l,2)
    ktrial(j,k,l,2)=kside-ktrial(j,k,la,2)+1
    if(l.eq.13) then
    w=ktrial(j,k,l,1)+k-1
    if(w.gt.kside) w=w-16
    ktrial(j,k,l,2)=w
    end if
50 continue
    do 60 l=14,16
    la=kside-l+2
    do 60 k=1,kside
    do 60 j=1,kside
    ktrial(j,k,l,2)=ktrial(j,k,l,2)
    ktrial(j,k,l,1)=ktrial(kside-j+1,k,la,1)
60 continue
c
c checkout coding follows
c
c    do 100 k=1,16
c    write(53,73)
c    write(53,73) k
c 73 format('set',i5)
c    write(53,73)
c    do 100 i=1,16
c    write(53,77) ((ktrial(i,j,k,l),j=1,16),l=1,2)
c 77 format(16i3,5x,16i3)
c 100 continue
c    close(53)
c    call exit
c    return
c    end

```

INSERT(jc,kc,lc,jd,kd,ld,apot,ltime,lz)

INSERT searches the list of afferent synapses on the desired neuron looking for a synapse with specific properties. If we find a match, we increment its potentiation by the value of apot; otherwise we establish a new synapse. The routine returns control to the caller.

```

subroutine insert(jc,kc,lc,jd,kd,ld,apot,ltime,lz)
include 'brain'
logical g,h
c
c jc    row of efferent neuron
c kc    column of efferent neuron
c lc    efferent nucleus
c jd    row of afferent neuron
c kd    column of afferent neuron
c ld    afferent nucleus
c apot  potential of synapse to be established
c ltime delay time of synapse
c lz=1  excitatory ipsilateral
c  =2   inhibitory ipsilateral

```

```

c  =3  excitatory contralateral
c  =4  inhibitory contralateral
c
      la=ksynct(jd,kd,ld)
c
c connection of up to 30 neurons allowed.
c multiple synapses are aggregated.
c
      if(la.le.0) go to 15
      do 10 kq=1,la
      g=ksynps(1,kq,jd,kd,ld).eq.jc
      g=g.and.(ksynps(2,kq,jd,kd,ld).eq.kc)
      g=g.and.(ksynps(3,kq,jd,kd,ld).eq.lc)
      g=g.and.(ksynps(4,kq,jd,kd,ld).eq.lz)
      g=(g.and.(ktime(kq,jd,kd,ld).eq.ltime))
      q=potent(kq,jd,kd,ld)
      if((lz.eq.1).or.(lz.eq.3)) then
      iop=1
      else
      iop=2
      end if
      if(lz.gt.2) apot=apot/4.0
      h=((iop.eq.1.and.q.ge.0.0).or.(iop.eq.2.and.q.le.0.0))
      if(g.and.h) then
      potent(kq,jd,kd,ld)=q+apot
      return
      end if
10  continue
15  continue
      if(la.ge.30) then
      jd=0
      kd=0
      ld=0
      return
      end if
      la=la+1
      ksynct(jd,kd,ld)=la
      ksynps(1,la,jd,kd,ld)=jc
      ksynps(2,la,jd,kd,ld)=kc
      ksynps(3,la,jd,kd,ld)=lc
      ksynps(4,la,jd,kd,ld)=lz
      potent(la,jd,kd,ld)=apot
      ktime(la,jd,kd,ld)=ltime
      synred(la,jd,kd,ld)=1.0
      return
      end

```

VISIIN

The next call by SETUP is to VISIIN an entry in VISUAL. VISIIN describes the division of visual space. The space extends 360 degrees horizontally, and from -20 to +50 vertically. We divide it into 256 regions for each eye.

```

subroutine visual
save
include 'brain'
dimension base(3,3),horiz(2,16),vert(2,16),ho(17),ve(17),
1 wall(3,4),aa(3),bb(3),c(3),d(3),e(3),f(3),g(3)
logical lck

```

```

a='l lateral geniculate      '
b='r lateral geniculate      '
call ident(a,jx,kl)
call ident(b,jx,kr)
call space(base)
high=0.5

```

```

do 5 j=1,kside
do 5 k=1,kside
histry(1,j,k,kl)=0.0
histry(1,j,k,kr)=0.0
5 continue

```

```

do 200 i=1,nrwall

```

```

do 10 k=1,3
wall(k,1)=awalls(k,1,i)
wall(k,2)=awalls(k,1,i)
wall(k,3)=awalls(k,2,i)
wall(k,4)=awalls(k,2,i)

```

```

10 continue
do 20 k=2,3
wall(2,k)=wall(2,k)+high

```

```

20 continue
do 25 l=1,3
f(l)=0.5*(body(1,l,knrrad-1)+body(2,l,knrrad-1))
aa(l)=wall(1,2)-wall(1,1)
bb(l)=wall(1,4)-wall(1,1)
d(l)=wall(1,1)

```

```

25 continue
call cross(aa,bb,c)
do 50 j=1,kside
do 50 k=1,kside
do 50 m=1,2
g(1)=horiz(2,k)*vert(2,16-j)
g(2)=vert(1,16-j)
g(3)=horiz(1,k)*vert(2,16-j)

```

```

c
c Whatever the meaning of left and right
c this will reverse the action.

```

```

c
if(m.eq.2) g(3)=-g(3)
do 30 ja=1,3
e(ja)=0.0
do 30 ka=1,3
e(ja)=e(ja)+base(ka,ja)*g(ka)

```

```

30 continue
   call veccos(c,e,wans)
   if(abs(wans).lt.0.01745) then
     lck=.false.
   else
     call punctr(c,d,e,f,g)
     do 35 kw=1,3
       zq=(g(kw)-f(kw))*e(kw)
       if(zq.lt.-0.000001) then
         lck=.false.
       go to 37
     end if
35 continue
   call inout(g,4,wall.lck)
   end if
37 continue
   if(lck) then
     call dstnce(f,g,ans)
     if(ans.gt.0.01) then
       dans=1.0/ans
     else
       dans=100.0
     end if
     if(m.eq.1) then
       if(histry(1,j,k,kl).lt.dans) histry(1,j,k,kl)=dans
     else
       if(histry(1,j,k,kr).lt.dans) histry(1,j,k,kr)=dans
     end if
   end if
50 continue

200 continue

   return

   entry visiin
   ho(1)=0
   ho(2)=1
   ho(3)=2
   ho(4)=3
   ho(5)=4
   ho(6)=5
   ho(7)=7
   ho(8)=9
   ho(9)=11
   ho(10)=15
   ho(11)=25
   ho(12)=40
   ho(13)=68
   ho(14)=96
   ho(15)=124
   ho(16)=152

```

```
ho(17)=180
```

```
ve(1)=-20
```

```
ve(2)=-10
```

```
ve(3)=-5
```

```
ve(4)=-4
```

```
ve(5)=-3
```

```
ve(6)=-2
```

```
ve(7)=-1
```

```
ve(8)=0
```

```
ve(9)=1
```

```
ve(10)=2
```

```
ve(11)=3
```

```
ve(12)=4
```

```
ve(13)=5
```

```
ve(14)=7
```

```
ve(15)=10
```

```
ve(16)=20
```

```
ve(17)=50
```

```
rads=6.28319/360.0
```

```
do 530 k=1,kside
```

```
hoz=rads*(ho(k+1)+ho(k))/2.0
```

```
vez=rads*(ve(k+1)+ve(k))/2.0
```

```
horiz(1,k)=sin(hoz)
```

```
horiz(2,k)=cos(hoz)
```

```
vert(1,k)=sin(vez)
```

```
vert(2,k)=cos(vez)
```

```
530 continue
```

```
return
```

```
end
```

MOTRIN

The last call by setup is to MOTRIN an entry in MOTR that sets parameters about the response of Pacrat to frustration.

We have now completed all the action under SETUP and return control to MAIN. MAIN calls LISTER.

```
subroutine motr
```

```
save
```

```
include 'brain'
```

```
ivade=ivade-1
```

```
if(ivade.lt.0) ivade=0
```

```
icont=icont-1
```

```
if(icont.lt.0) icont=0
```

```
10 if(ivade.gt.0) then
```

```
call newdir(kdirec)
```

```
go to 215
```

```
end if
```

c

```

a='l motor turnout
call ident(a,j,k)
bigl=0.0
do 200 ka=1,kside
do 200 kb=1,kside
if(histry(1,ka,kb,k).gt.bigl) bigl=histry(1,ka,kb,k)
200 continue
a='r motor turnout
call ident(a,j,k)
bigr=0.0
do 210 ka=1,kside
do 210 kb=1,kside
if(histry(1,ka,kb,k).gt.bigr) bigr=histry(1,ka,kb,k)
210 continue
if((bigl.ge.atriga).and.(bigr.ge.atriga))then
ivade=70
if(rand(x).gt.0.5) then
kdirec=3
else
kdirec=4
end if
go to 10
end if
if((bigl.ge.atrig).or.(bigr.ge.atrig))then
icont=5+2*rand(x)
if(bigl.ge.bigr) then
call newdir(3)
write(38,*) icnt,'tactil went left'
go to 215
else
call newdir(4)
write(38,*) icnt,'tactil went right'
go to 215
end if
else
if(ivade.gt.0) go to 213
wa=videol(x)
wb=videor(x)
if((wa+wb).gt.0.0) then
if(icont.gt.0) go to 213
if(wa.ge.wb) then
call newdir(4)
write(38,*) icnt,'vision went right'
go to 215
else
call newdir(3)
write(38,*) icnt,'vision went left'
go to 215
end if
end if
end if

```

```

213 continue
    dxm=dxm*1.05
    if(dxm.gt.0.05) dxm=0.05
    go to 220
215 if(dxm.gt.0.05) then
    dxm=dxm*0.9
    else
    dxm=dxm*0.95
    end if
    if(dxm.lt.0.02) dxm=0.02
c
220 continue
    kckm=0
222 continue
    avectr(1)=dxm
    phase=phase+5.0*dxm
    if(phase.gt.6.28319) phase=phase-6.28319
    call movpac
    if((dxm.lt.0.0).and.(kckm.eq.0))then
    kckm=1
    go to 222
    end if
c
    if(mod(icnt,10).eq.0) then
    write(4,240) icnt, knrrad
    240 format(2i10)
    do 250 k=1, knrrad
    do 250 i=1,2
    write(4,242) (body(i,j,k),j=1,3)
242 format(3f10.5)
250 continue
    end if
    return
    entry motrin
    ivade=0
    atrig=0.001
    atriga=atrig*20.0
    return
    end

```

LISTER

LISTER is a last step in preparing the brain to run. It writes out the names of the centers and of the nuclei. Then it makes a complete analysis of the net and also writes it to the file. It tabulates the efferent and afferent synapses of each nucleus. Following this recapitulation, we return control to MAIN for the last time before execution proper.

```

subroutine lister
include 'brain'
dimension ktemp(1000,3,2)
write(30,*) 'centers'
write(30,*)

```

```

do 5 k=1,nrctr
5  write(30,*) k, ' ',centrs(k)
   write(30,*)
   write(30,*) 'nuclei'
   write(30,*)
   do 10 k=1,nrnucs
10  write(30,*) k, ' ',nuclei(k)
   do 20 l=1,nrnucs
   if(kruse(l).eq.0) go to 20
   write(18,15) nuclei(l),
1   alertd(l),runavg(l)
15  format(1h ,a30,5x,3f10.5)
20  continue
   do 100 mx=1,nrctr
   write(18,35) centrs(mx)
35  format(1h0,' *** ',a30)
   do 90 l=kcnuks(mx,1),kcnuks(mx,2)
   if(kruse(l).eq.0) go to 90
   write(18,55) nuclei(l)
55  format(1h0,a30)
   write(18,57)
57  format(1h ,5x,'genesis')
   if(krnet(l).eq.0) then
   write(18,59)
59  format(1h ,16x,'nowhere')
   else
   do 70 kr=1,krnet(l)
   write(18,63) nucnet(l,kr,2),nuclei(nucnet(l,kr,1))
63  format(1h ,10x,i5,x,2a30)
70  continue
   end if
   write(18,65)
65  format(1h ,5x,'epigenesis')
   if(krgro(l).eq.0) then
   write(18,59)
   else
   do 80 kg=1,krgro(l)
   write(18,63) nucgro(l,kg,2),nuclei(nucgro(l,kg,1))
80  continue
   end if
   write(18,82)
82  format(1h ,5x,'genetic afferents from')
   kvinc=0
   do 85 la=1,nrnucs
   if(krnet(la).eq.0) go to 85
   do 84 lax=1,krnet(la)
   if(nucnet(la,lax,1).ne.1) go to 84
   kvinc=1
   write(18,63) nucnet(la,lax,2),nuclei(la)
84  continue
85  continue
   write(18,86)

```



```

86 format(1h ,5x,'epigenetic afferents from')
   do 88 la=1,nrnucs
   if(krgro(la).eq.0) go to 88
   do 87 lax=1,krgro(la)
   if(nucgro(la,lax,1).ne.1) go to 87
   kvinc=1
   write(18,63) nucgro(la,lax,2),nuclei(la)
87 continue
88 continue
   if(kvinc.eq.0) write(18,59)
90 continue
100 continue
   do 110 l=1,nrnucs
   do 110 ka=1,3
   do 110 la=1,2
110 ktemp(ja,ka,la)=0
   do 150 l=1,nrnucs
   if(kruse(l).eq.0) go to 150
   do 125 k=1,kside
   do 125 j=1,kside
   if(ksynct(j,k,l).eq.0) go to 125
   do 120 i=1,ksynct(j,k,l)
   ja=ksynps(1,i,j,k,l)
   ka=ksynps(2,i,j,k,l)
   la=ksynps(3,i,j,k,l)
   if(potent(i,j,k,l).ge.0.0) then
   ktemp(l,1,1)=ktemp(l,1,1)+1
   ktemp(la,2,1)=ktemp(la,2,1)+1
   if(l.eq.la) then
   ktemp(l,3,1)=ktemp(l,3,1)+1
   end if
   else
   ktemp(l,1,2)=ktemp(l,1,2)+1
   ktemp(la,2,2)=ktemp(la,2,2)+1
   if(l.eq.la) then
   ktemp(l,3,2)=ktemp(l,3,2)+1
   end if
   end if
120 continue
125 continue
150 continue
   do 190 l=1,nrnucs
   if(kruse(l).eq.0) go to 190
   write(18,185) nuclei(l),(ktemp(l,jn,1),jn=1,3)
185 format(1h ,a30,' afferents ',i5,' efferents ',
   1 i5,' self ',i5)
   write(18,187) (ktemp(l,jn,2),jn=1,3)
187 format(1h ,5x,'inhibiting',26x,i5,11x,i5,6x,i5)
190 continue
   return
end

```

The Main Loop

We have built the brain, the organism, and the universe. The life of Pacrat begins.

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APPENDIX

character*30 a,b,centrs,nuclei,nuc1st
logical chkout,hotstr,flags

common/names/ nrctrs,nrnucs,centrs(100),nuclei(1000),
1 kcnucs(100,2),kruse(1000),krnet(1000),krgro(1000),
2 nucnet(1000,20,4),nucgro(1000,20,4),
3 kcensz,knucsz,knucwd,stinc,knucs,kside

common/abrain/
1 ksynps(4,30,16,16,300),potent(30,16,16,300),
2 synred(30,16,16,300),ksynct(16,16,300),
3 histry(10,16,16,300),ktime(30,16,16,300)

common/dynam/ alertd(1000), runavg(1000), thresh(16,16,300)

common/plastk/ recptv(16,16,300),grfctr(16,16,300),
1 synare(2,2,16,16,300),grinc

common/updatp/ reduce,frustr,frivla,frivlb,
1 rewavg,recovr,thrup,thrdown,whz

common/run/ chkout,hotstr,ksnap,flags(100),nuc1st(100),
1 kbndls(3),lstnr(10),icnt,kbgend,klend

common/pacrt/ knrrad,radii(2,3,100),body(2,3,100),nrwall,
1 awalls(3,2,100)

common/movng/ avectr(3),amoton(3,3),bmoton(3,3,100),
1 amovr(3,100),phase,radi(2,3,100)

common/transf/ snpass(16,16,100),ampass(16,16,100),dist,
1 disr(2),dxm

dimension alatgl(16,16),alatgr(16,16),venpl(16,16),
1 venpr(16,16),area4l(16,16),area4r(16,16)
equivalence (alatgl,snpass),(alatgr(1),snpass(1,1,2)),
1 (venpl(1),snpass(1,1,3)),
2 (venpr(1),snpass(1,1,4)),
3 (area4l,ampass(1,1,3)),
4 (area4r(1),ampass(1,1,4))
a

a character variable available for general
use.

alertd(1000) dynam
level of general activation of a nucleus.

amoton(3,3) movng
a movement matrix.

amovr(3,100) **movng**
a movement matrix.

ampass(16,16,100) **transf**
used to pass motor information from Pacrat
to the universe.

avectr(3) **movng**
displacement of the center of Pacrat's nose.

awalls(3,2,100) **pacrt**
specifications of the walls.

b
a character variable available for general
use.

bmoton(3,3,100) **movng**
a movement matrix.

body(2,3,100) **pacrt**
actual position of Pacrat's body.

centrs(100) **names**
names of the brain centers.

chkout **run**
flag that determines whether intermediate
listings are desired.

disr(2) **transf**
temporary variables used in tactil for
distance check.

dist **transf**
minimum distance from Pacrat to the closest
wall.

dxm **transf**
the basic increment of movement.

flags(100) **run**
the flags that determine the combination of

intermediate printouts.

frivla updatp
the upper frustration level check.

frivlb updatp
the lower frustration level check.

frustr updatp
the frustration level (an emotion).

grfctr(16,16,300) plastk
the pressure on an axon to grow.

grinc plastk
the potentiation of a new synapse.

histry(10,16,16,300)abrain
activation history of a neuron.

hotstr run
flag for a hot start.

icnt run
counter for time steps.

kbgend run
the outer loop in the main program.

kbndls(3) run
three parameters for saving histories
(equivalent to do loop.)

kcensz names
upper limit on number of centers.

kcnucs(100,2) names
first and last nucleus in each center.

kltend run
inner loop in main program.

knrrad pacrt
number of body segments.

knucs names
total number of nuclei.

knucsz names
limit on number of nuclei.

knucwd names
upper limit on number of nuclei this nucleus
may be efferent upon - or grow to.

kgro(1000) names
the actual number of nuclei this nucleus may
grow to.

krnet(1000) names
the actual number of nuclei this nucleus is
genetically efferent upon.

kruse(1000) names
type of nucleus (sensor, motor, other).

kside names
the number of neurons on one side of a
nucleus (they are square.)

ksnap run
interval for snapshots (mod(icnt,ksnap))

ksynct(16,16,300) abrain
count of synapses on this neuron.

ksynps(4,30,16,16,3)abrain
description of afferent neuron: 1 row, 2
column, 3 nucleus, 4 type.

ktime(30,16,16,300)abrain
axonal delay to synapse.

lstnr(10) run
number of nuclei to be listed in this

printout.

nrctr names
the number of centers.

nrnucl names
the number of nuclei.

nrwall pacrt
the number of walls.

nucgro(1000,20,4) names
growth: 1 nucleus to, 2 type of
distribution, 3 type of neuron, 4 type of synapse.

nuclei(1000) names
names of nuclei.

nuc1st(100) run
names of nuclei to be listed by CHEKPR.

nucnet(1000,20,4) names
genetic: 1 nucleus to, 2 type of
distribution, 3 type of neuron, 4 type of synapse.

phase movng
the phase of the sinusoidal movement of
Pacrat.

potent(30,16,16,300)abrain
potentiation of a synapse.

radi(2,3,100) movng
working position of Pacrat.

radii(2,3,100) pacrt
absolute position of Pacrat.

recovr updatp
increment of dynamic synaptic condition.

recptv(16,16,300) plastk
receptivity of a neuron to the formation of
synapses.

reduce updatp
decrement of dynamic synaptic condition.

rewavg updatp
used in coding for inhibitory growth.

runavg(1000) dynam
running average of nuclear activity.

snpass(16,16,100) transf
used to pass sensory information from the
universe to Pacrat.

stinc names
original potentiation of a genetic synapse.

synare(2,2,16,16,30)plastk
synaptic area: from this nucleus and from
others.

synred(30,16,16,300)abrain
readiness of this synapse.

thrdwn updatp
hypopolarization increment.

thresh(16,16,300) dynam
dynamic threshold of this neuron.

thrup updatp
hyperpolarization increment.

whz updatp
used in hillock and in normal.

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